

## **CERTIFICATE FOR INCLUSION OF A MEDICAL DEVICE**

**ARTG Number** 137139

**ARTG Labelname** Shanree Pty Ltd T/A Epi-Assist Monitoring - Alarm, epilepsy

**Sponsor** Shanree Pty Ltd T/A Epi-Assist Monitoring

**Commencement Date** 03/04/2007

**Manufacturer** Emfit Ltd Finland

**Device Class** Class 1

**GMDN Code** 36693 Alarm, epilepsy

**ARTG Product Number and Name**  
223850 Alarm, epilepsy

**The above Medical Device is Included in the Australian Register of Therapeutic Goods subject to the following conditions**

***Standard Conditions***

*The automatic conditions applicable to the inclusion of all kinds of medical devices in the Register are as specified in section 41FN of the Therapeutic Goods Act 1989.*

*The standard conditions that are imposed under section 41FO of the Therapeutic Goods Act 1989 when kinds of medical devices are included in the Register are as set out in the following paragraphs.*

*For a medical device included in the Register under Chapter 4 and imported into Australia, the Sponsor must ensure that information about the Sponsor is provided in such a way as to allow the sponsor to be identified.*

*Each sponsor shall retain records of the distribution of all of the sponsor's medical*

*devices included in the Register under Chapter 4. In the case of records relating to a Class AIMD medical device, Class III medical device, or Class IIb medical device that is an implantable medical device, the distribution records shall be retained for a minimum period of 10 years. In the case of records relating to any other device, the distribution records shall be retained for a minimum period of 5 years.*

*The sponsor of a medical device included in the Register under Chapter 4 shall keep an up to date log of information of the kind specified in Regulation 5.8.*

*The sponsor shall provide to the Director, Office of Devices, Blood and Tissues, Therapeutic Goods Administration, three consecutive reports which include information of a kind specified in Regulation 5.8 that arises during the reporting period, concerning Class III, Class AIMD or implantable Class IIb medical devices. The reporting period for the first report commences on the date of inclusion and is to be at least a period of six months but not more than 18 months ending on 1 October. For subsequent reports the reporting period is 12 months ending on 1 October.*

*Where a medical device included in the Register, contains a substance which is included in the Fourth Schedule to the Customs (Prohibited Imports) Regulations or the Eighth Schedule to the Customs (Prohibited Exports) Regulations the Sponsor shall, at the time of importation or exportation of the medical device, be in possession of a licence and a permission for importation or exportation of each consignment of the goods as required by those regulations.*

*A sponsor shall ensure that a medical device within their control is stored and transported in accordance with the instructions and information provided by the manufacturer.*